

# Adapting the Burrows-Wheeler Transform (BWT) for Personalized DNA Analysis

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CS460



# Introduction

- **Objective:** Adapt the BWT algorithm to enhance personalized DNA analysis.
- **Importance:** Empower users with private, efficient genetic data insights.
- **BWT Introduction:** Overview of BWT's role in data compression and pattern matching.

# Project Overview

**Problem:** Efficient, private analysis of raw DNA data from services like 23andMe.

**Solution:** Use BWT for data compression and analysis on personal computers.

**Benefits:**

- Increased accessibility of DNA analysis.

- Ensures user privacy by performing analysis locally.

- Improved efficiency in searching genetic data.

# Methodology

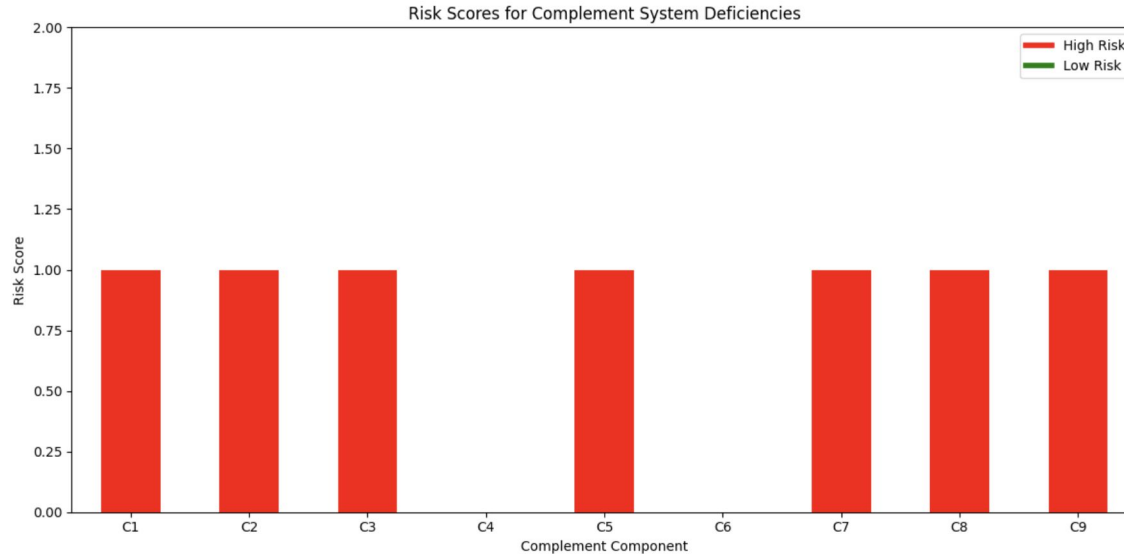
## Content:

- **Data Processing:**
  - Reading raw DNA data files.
  - Extracting relevant genetic information (rsid, chromosome, position, genotype).
- **BWT Application:**
  - Transforming genetic data string using BWT.
  - Steps in BWT encoding and decoding.

```
1 def read_23andme(file_path):
2     # Open the file at the specified path in read mode
3     with open(file_path, 'r') as file:
4         # Read all lines from the file
5         lines = file.readlines()
6
7     # Filter out lines that start with '#' (comments) and strip any leading/trailing whitespace
8     data_lines = [line.strip() for line in lines if not line.startswith('#')]
9
10    # Split each line by tab characters to separate the columns
11    data = [line.split('\t') for line in data_lines]
12
13    # Create a DataFrame from the processed data with appropriate column names
14    df = pd.DataFrame(data, columns=['rsid', 'chromosome', 'position', 'genotype'])
15
16    # Return the DataFrame
17    return df
```

# Risk Assessment for Complement Deficiencies

	rsid	chromosome	position	genotype
0	rs548049170	1	69869	TT
1	rs9283150	1	565508	AA
2	rs116587930	1	727841	GG
3	rs3131972	1	752721	GG
4	rs12184325	1	754105	CC



# Results

## Content:

- **BWT Transformation:**
  - Example output of BWT transformation.

main.py

```
1 genetic_string = ''.join(df['rsid'] + df['genotype'])
2 bwt_result = bwt_transform(genetic_string)
```

## Genetic Risk Analysis:

- AMD and Breast Cancer risk scores.
- Example risk score calculation.

```
_snps = {'rs1061170': 'C', 'rs3753394': 'A', 'rs10490924': 'T', 'rs2230199': 'G', 'rs9332739': 'A'}
data = df[df['rsid'].isin(_snps.keys())].copy()
```

# Conclusion and Future Work

## **Summary:**

- Successful adaptation of BWT for DNA analysis.
- Enhanced privacy and efficiency.

## **Future Enhancements:**

- Expanding analysis to include more genetic markers.
- Integration with other health data for comprehensive analysis.